REPORT DOCUMENTATION PAGE Form Approved OMB NO. 0704-0188 The public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. regarding this burden estimate or any other aspect of this collection of information, including suggesstions for reducing this burden, to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204, Arlington VA, 22202-4302. Respondents should be aware that notwithstanding any other provision of law, no person shall be subject to any oenalty for failing to comply with a collection of information if it does not display a currently valid OMB control number. PLEASE DO NOT RETURN YOUR FORM TO THE ABOVE ADDRESS. 1. REPORT DATE (DD-MM-YYYY) 2. REPORT TYPE 3. DATES COVERED (From - To) 29-02-2012 Abstract 4. TITLE AND SUBTITLE 5a. CONTRACT NUMBER A Novel Methodology for Exposing Tissue Cultures to Blast W911NF-10-1-0526 Overpressure for Determining Injury Criteria 5b. GRANT NUMBER 5c. PROGRAM ELEMENT NUMBER 611103 6. AUTHORS 5d. PROJECT NUMBER Matthew B. Panzer, Cameron R. 'Dale' Bass, Bruce P. Capehart, Barclay Morrison III, David F. Meaney 5e. TASK NUMBER 5f. WORK UNIT NUMBER 7. PERFORMING ORGANIZATION NAMES AND ADDRESSES 8. PERFORMING ORGANIZATION REPORT NUMBER University of Pennsylvania Office of Research Services University of Pennsylvania Philadelphia, PA 19104 -9. SPONSORING/MONITORING AGENCY NAME(S) AND 10. SPONSOR/MONITOR'S ACRONYM(S) ADDRESS(ES) ARO 11. SPONSOR/MONITOR'S REPORT U.S. Army Research Office NUMBER(S) P.O. Box 12211 Research Triangle Park, NC 27709-2211 58155-LS-MUR.10 12. DISTRIBUTION AVAILIBILITY STATEMENT Approved for public release; distribution is unlimited. 13. SUPPLEMENTARY NOTES The views, opinions and/or findings contained in this report are those of the author(s) and should not contrued as an official Department of the Army position, policy or decision, unless so designated by other documentation. 14. ABSTRACT Introduction: A growing number of clinicians and scientists acknowledge that there exists some level of primary blast exposure that can disrupt or damage neuronal tissue resulting in traumatic brain injury. This injury may be associated with persistent and progressive sequelae. While the groundwork for identifying the brain's tolerance to primary blast has been created using animal and computational models, the mode (volumetric or deviatoric) and threshold of tissue-level deformation associated with brain tissue injury remains unknown. We report here a new 15. SUBJECT TERMS

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A Novel Methodology for Exposing Tissue Cultures to Blast Overpressure for Determining Injury Criteria

ABSTRACT

Introduction: A growing number of clinicians and scientists acknowledge that there exists some level of primary blast exposure that can disrupt or damage neuronal tissue resulting in traumatic brain injury. This injury may be associated with persistent and progressive sequelae. While the groundwork for identifying the brain's tolerance to primary blast has been created using animal and computational models, the mode (volumetric or deviatoric) and threshold of tissue-level deformation associated with brain tissue injury remains unknown. We report here a new test methodology to apply overpressure conditions to brain tissue preparations in vitro similar to those created from blast. We hypothesize this method would replicate the physical characteristics of blast, allowing measurements of tissue mechanical response and ultimately cell and tissue injury response.

Materials and Methods: Blast waves were generated by a helium-driven shock tube (76 mm diameter tube with a 25 mm driver and a 1220 mm driven section). The blast wave impacts a fluid-filled apparatus that contains living brain tissue, creating a single overpressure pulse that propagates through the suspended tissue. Downstream, the apparatus was designed to mitigate returning shock reflections. An axisymmetric numerical model of the apparatus and tissue was developed in LS-DYNA to assess the response of the tissue that could not be measured experimentally. Experimental and modeled results were compared.

Results and Discussion: The apparatus was able to generate blast overpressure waves at the tissue culture that were characteristic of Friedlander blast waves with minimal amount of reflected overpressure, but with slightly longer rise times (Figure 1A). Peak reflected overpressures generated using the shock tube ranged from 500 to 1500 kPa with durations from 0.5 to 1.0 ms. The numerical model was validated using the experimental results for both the external shock tube overpressure response and the fluid-filled environment at the location of the tissue culture (Figure 1B). Analysis of the numerical model showed minimal deviatoric deformation in the tissue culture.

Conclusions: The test apparatus was developed to simulate the in vivo conditions of tissue within a brain exposed to a single blast wave by sustaining the tissue culture in fluid that has been impacted by a shock tube-generated blast. Since results indicate that the experimental apparatus was effective in isolating the tissue from deviatoric stress, this apparatus and model can be utilized for studying overpressure-based injury criteria. Pressure-based injury criteria will be further enhanced in future studies by extending the range of input overpressures on the living brain tissue, as well as identifying additional measures of histopathologic tissue damage and functional impairment from electrophysiology measurements.

A Novel Methodology for Exposing Tissue Cultures to Blast Overpressure for Determining Injury Criteria

Target Track: Neural Engineering

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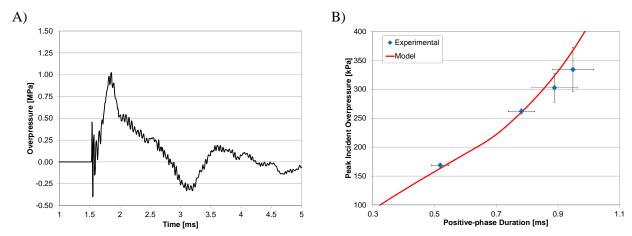


Figure 1: A) A typical overpressure time-history at the tissue produced by the shock tube, and B) the validation of the shock tube model using the external shock tube overpressure response

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